



Docket No. 20784/6

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

APPLICANT: Hans-Ulrich Demuth et al.

ART UNIT: 1646

SERIAL NO.: 10/082,001

FILED:

February 22, 2002

FOR: NEW EFFECTORS OF DIPEPTIDYL PEPTIDASE IV FOR TOPICAL USE

CERTIFICATE OF MAILING I hereby certify that this paper (along with any paper referred to as being attached or enclosed) is being deposited with the United States Postal Service on the date shown below with sufficient postage as first class mail in an envelope addressed to the: Commissioner for Patents, Washington, D.C., 20231-0001 October 28, 2002 Sandra J. Graves

INFORMATION DISCLOSURE STATEMENT

Commissioner for Patents Washington, D.C. 20231

Date: October 28, 2002

Pursuant to Applicant(s) duty of disclosure, the information listed in the attached form PTO-1449 is brought to the attention of the Examiner. Copies of the listed items are enclosed.

The citation of the listed items is not a representation that they constitute a complete or exhaustive listing of the relevant art or that the references are prior art. The items listed are submitted in good faith, but are not intended to substitute for the Examiner's search. It is hoped, however, that in addition to apprising the Examiner of these particular items, they will assist in identifying fields of search and in making as full and complete a search as possible.

The filing of this information disclosure statement is not an admission that the information cited herein is, or is considered to be, material to patentability as defined in 37 C.F.R. § 1.56(b).

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In accordance with the requirement under 37 C.F.R. 1.98 (3)(i), the following are concise explanations of the relevance, as presently understood, with regard to those items submitted herewith that are not in the English language:

I. <u>LIST OF PATENTS, PUBLICATIONS OR OTHER INFORMATION</u>

The patents, publications or other information submitted for consideration by the Office are listed on PTO-1449, attached hereto.

II. COPIES

a. Submitted herewith is a legible copy of (i) each U.S and foreign patent; (ii) each publication or that portion which caused it to be listed; and (iii) all other information or that portion which caused it to be listed

III. CONCISE EXPLANATION OF THE RELEVANCE

(check at least one box)

- a. Except as may be indicated below in (b), all of the patents, publications or other information are in the English language or were cited in an English language Search Report, a copy of which is attached hereto (concise explanation not required).
- b. A concise explanation of the relevance of all patents, publications or other information listed that is not in the English language is as follows:

FRENCH LANGUAGE FR 2 696 740 Applicants have relied on an English language abstract in determining that this patent apparently relates to Dimethylbiguanide drug derivatives and their medical applications.

FRENCH LANGUAGE FR 2 085 665 Applicants have relied on an English language abstract in determining that this patent apparently relates to biguanide substitutes possessing a hypoglycemic property.

GERMAN LANGUAGE DT 25 42 598 A1 Applicants have relied on an English language abstract in determining that this patent apparently relates to biguanide salts as well as a method for preparing such salts. In addition to this the invention apparently covers pharmaceutical compounds which contain such salts.

GERMAN LANGUAGE WO 97/40832 Applicants have relied on an English language abstract in determining that this patent apparently relates

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to the use of a method for reducing in the blood of a mammal by administration of effectors the enzyme activity of dipeptidyl peptidase (DP IV) or enzyme activity similar to DP IV, the endogenous (or additionally exogenously administered) insulinotropic peptide gastric inhibitory polypeptide 1-42 and glucagon-like peptide amide-1 7-36 are decomposed in a causal sequence to a reduced extent by DP IV enzymes or those similar to DP IV.

JAPANESE PATENT JP 4-334357A2: Applicants have relied on an English language abstract in determining that this patent ACYL DERIVATIVE HAVING ENZYME-INHIBITING ACTION, apparently relates to a compound having a prolyl endopeptibase activity-inhibiting action and useful as an antidement agent, especially an anti-amnestic agent.

GERMAN LANGUAGE DE 299 09 210 U1 Applicants have relied on an English language abstract in determining that this patent apparently relates to dipeptide compounds or compounds analogous to dipeptide compounds, which are made of an amino acid and a thiazolidine or pyrrolidine group, and to their salts. The invention further relates to the use of these compounds in the treatment of impaired glucose tolerance, glucosuria, hyperlipidemia, metabolic acidosis, diabetes mellitus, diabetic neuropathy and nephropathy as well as secondary diseases of diabetes mellitus in mammals.

c. The following additional information is provided for the Examiner's consideration:

FEES

IV. THIS IDS IS BEING FILED UNDER 37 C.F.R. § 1.97(b) (check one box)

a. 🔲	within three months of the filing date of a national application (37	C.F.R.
	§ 1.97(b) (1)). No fee or certification is required.	

- b. within three months of the date of entry of the national stage as set forth in §1.491 in an international application (37 C.F.R. § 1.97(b) (2)). No fee or certification is required.
- c. (3) before the mailing date of a first Action on the merits (37 C.F.R. § 1.97(b) (3)). No fee or certification is required. In the event that a first Office Action on the merits has been issued, please consider this IDS under 37 C.F.R. § 1.97(c) and see the certification under 37 C.F.R. § 1.97(e) below,

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or, if no certification has been made, charge our deposit account a fee in the amount of \$180.00 as required by 37 C.F.R. § 1.17(p).

V.		IDS IS BEING FILED UNDER 37 C.F.R. § 1.97(c): c one box)
	1.97(c	the mailing date of a Final Office Action under 37 C.F.R. § 1.113 (See 37 C.F.R. §) (1)) or before the mailing date of a Notice of Allowance under 37 C.F.R. § 1.311 7 C.F.R. § 1.97(c) (2)).
	a. 🗌	No certification; therefore, a fee in the amount of \$180.00 is required by 37 C.F.R. § 1.17(p).
	b. [_	or See the certification below. No fee is required.
VI.	CERT	IFICATION UNDER 37 C.F.R. § 1.97(e) (check only one box)
The ur	ndersign	ned hereby certifies that
	a. 🗌	each item of information contained in the IDS was cited in a communication from a foreign Patent Office in a counterpart foreign application not more than three months prior to the filing of this IDS; or
	b. 🗌	no item of information contained in the IDS was cited in a communication from a foreign Patent Office in a counterpart foreign application or, to the best of my knowledge after making reasonable inquiry, was known to any individual designated in 37 C.F.R. § 1.56(c) more than three months prior to the filing of this statement.
	c	Some of the items of information were cited in a communication from a foreign Patent Office as indicated in the Form 1449 by those references having an asterisk(*). As to this information, the undersigned certifies that each item of information contained in the IDS was cited in a communication from a foreign Patent Office in a counterpart foreign application not more than three months prior to the filing of this IDS. As to the remaining information, the undersigned hereby certifies that no item of this remaining information contained in the IDS was cited in a communication from a foreign Patent Office in a counterpart foreign application or, to the best of my knowledge after making reasonable inquiry, was known to any individual designated in 37 C.F.R. § 1.56(c) more than three months prior to the filing of this statement.
		A check in the amount of \$180.00 is enclosed for the above-indicated fee. A duplicate copy of this paper is attached.

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\boxtimes	No f	ee is required.
VII.	THIS IDS IS	S BEING FILED UNDER 37 C.F.R. § 1.704(d) (PATENT TERM
	<u>ADJUSTMI</u>	ENT)
	Applies to o	original applications (other than design) filed on or after May 29, 2000.
a.		Each item of information contained in the Information Disclosure
		Statement was cited in a communication from a foreign patent office in a
		counterpart application and this communication was not received by any
		individual designated in § 1.56© more than thirty days prior to the filing
		of the Information Disclosure Statement.
b.	X	Enclosed herewith is form PTO-1449.
c.	<u>X</u>	Copies of cited references are enclosed.
d.		The listed references were cited in the enclosed International Search

If the Examiner has any questions concerning this IDS, the Examiner is requested to contact the undersigned. If it is determined that this IDS has been filed under the wrong rule, the PTO is requested to consider this IDS under the proper rule (with a petition, if necessary) and charge any additional fees to Deposit Account No. 50-0369.

Report in a counterpart foreign application.

Date: October 28, 2002

Respectfully submitted,

John C. Serio

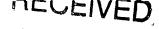
Reg. No. 39,023

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FORM PTO-1449 INFORMATION DISCLOSURE STATEMENT		ATTY. DOCKET NO. SERIAL NO. 10/082,001					
			APPLICANT(S): Demuth et al.				
				FILING DATE: February 22, 2002		ART UNIT	:
			UNITED STAT	ES PATENT DOCUMENTS	•		
EXAM. INITIAL		DOCUMENT NUMBER	DATE	INVENTOR	CLASS	SUB CLASS	FIL. DATE IF APPR
	AA	2,961,377	11/22/1960	Shapiro et al.	167	65	
	AB	3,174,901	03/23/1965	Sterne	167	65	
	AC	3,879,541	04/22/1975	Kabbe et al.	424	326	
	AD	3,960,949	06/01/1976	Ahrens et al.	260	564 B	
	AE	4,028,402	06/07/1977	Fischer et al.	260	501.14	
	AF	4,935,493	06/19/1990	Bachovchin et al.	530	331	
	AG	5,433,955	07/18/1995	Bredehorst et al.	424	94.3	
	AH	5,462,928	10/31/1995	Bachovchin et al.	514	19	
	ΑI	5,512,549	4/30/1996	Chen et al.	514	12	
	AJ	5,543,396	08/06/1996	Powers et al.	514	19	
	AK	5,614,379	03/25/1997	MacKellar	435	68.1	
	AL	5,624,894	04/29/1997	Bodor	514	2	
	AM	5,939,560	08/17/1999	Jenkins et al.	548	535	
	AN	6,006,753	12/28/1999	Efendic	128	898	
			FOREIGN P	ATENT DOCUMENTS	- u		
		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB CLASS	TRAN Y/N
	BA	WO 01/62266 A2	08/30/2001	PCT	A61K	38/00	Y
	ВВ	WO 00/53171	09/14/2000	PCT	A61K	31/155	Y
•	вс	DT 25 42 598 A1	04/22/1976	Germany	C07C	129/16	N
	BD	FR 2 696 740 A1	04/15/1994	France	C07D	207/404	N
	BE	FR 2 085 665	12/31/1971	France	A61K	27/00	N
	BF	WO 97/40832	11/06/1997	PCT	A61K	31/425	Abstract only
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ATTY. DOCKET GOOS THR 1600/290 PRIAL NO. TRADEMATORM PTO-1449 10/082,001 INFORMATION DISCLOSURE STATEMENT 20784/6 APPLICANT(S): Demuth et al. ART UNIT: FILING DATE: 1646 February 22, 2002 FOREIGN PATENT DOCUMENTS **CLASS SUB TRAN COUNTRY DOCUMENT** DATE **CLASS** Y/N NUMBER Y 207/16 WO 95/15309 06/08/1995 **PCT** C07D BG C07C 233/57 Abstract Japan BH JP 4334357 11/20/1992 only Y **PCT** C12N WO 93/08259 04/29/1993 BIA61K 37/00 Y **PCT** 05/04/1995 BJWO 95/11689 Y A61K 31/435 **PCT** BK WO 97/45117 12/04/1997 Y A61K 45/00 BLDE 196 16 486 C2 10/30/1997 Germany Y 38/00 **PCT** A61K WO 95/29691 11/09/1995 BM C07K 5/06 Y WO 98/22494 05/28/1998 **PCT** BN Y **PCT** C12Q 1/68 WO 00/01849 01/13/2000 BO 14/605 Y **EPO** C07K 06/21/1995 BP EP 0 658 568 A1 295/04 Y C07D BQ DD 296 075 A5 11/21/1991 Germany 295/04 N C07D BR DD 296 075 A5 11/21/1991 Germany Y 15/16 04/24/1996 **EPO** C12N BS EP 0 708 179 A2 31/425 N **EPO** A61K BT EP 0 995 440 A1 04/26/2000 7/34 Y 08/08/1991 **PCT** C07K BU WO 91/11457 C07K 5/10 Y **PCT** 10/31/1991 BV WO 91/16339 Y C07D 207/00 **PCT** BW WO 98/19998 05/14/1998 37/54 Y **PCT** A61K BXWO 91/17767 11/28/1991 (Abstract 10/13/1992 A61K 37/64 BY JP 04-288098 JP Only) Y 9/572 WO 99/46272 A 09/16/1999 **PCT** C07F BZA61K 31/425 N BAA DE 299 09 210 U 09/09/1999 Germany

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FORM PTO-1449 INFORMATION DISCLOSURE STATEMENT		ATTY. DOCKET NO. 20784/6	SERIAL NO. 10/082,001	
		APPLICANT(S): Demuth et al.		
		FILING DATE: February 22, 2002	ART UNIT: 1646	
	OTHER DOCUMENTS (INCLUDING A	UTHOR, TITLE, DATE, PERTINENT	Γ PAGES, ETC.)	
CA	Campbell, I.W. New Antidiabetic Drugs, edefficacy and inadequacy". 3:33-51 (1990).	l. C.J. Bailey & P.R. Flatt, Smith-Gordon,	"Sulphonylureas and metforming	
СВ	The Merck Index, 11 th Edition, An Encyclop	pedia of Chemicals, Drugs, and Biological	ls, 1989, Page 934	
CC	The Merck Index, 12 th Edition, An Encyclop	pedia of Chemicals, Drugs, and Biological	ls, 1996, Page 1014.	
CD	Martindale The Extra Pharmacopoeia, 30th E	Edition, London Pharmaceutical Press, 199	93, Page 1619.	
CE	Martindale The Extra Pharmacopoeia, 30th E	Edition, London Pharmaceutical Press, 199	93, Page 36.	
CF	CHEMICAL ABSTRACTS, vol. 115. No. 15, 14. October 1991 (1991-10-14) Columbus, Ohio, US; abstract no. 149947q, SCHOEN EKKEHARD ET AL: "Dipeptidyl peptidase IV in the immune system. Effects of specific enzyme inhibitors on activity of dipeptidyl peptidase IV and proliferation of human lymphocytes"			
CG	CHEMICAL ABSTRACTS, vol. 126, no. 2, 13. January 1997 (1997-01-13) Columbus, Ohio, US; abstract no. 16161j, STOECKEL A. ET AL: "Competitive inhibition of proline specific enzymes by amino acid thioxopyrrolidides and thiazolidides".			
СН	CHEMICAL ABSTRACTS, vol. 118, no. 25, 21. June 1993 (1993-06-21) Columbus, Ohio, US; abstract no. 255342k, Hosoda, et al, "Preparation of N-(heterocyclic Carbonyl) Amino Acids and Analogs as Prolyl Endopeptidase Inhibitors", November 1992 (1992-11-20)			
CI	ARAI ET AL., "Synthesis of prolyl endopep vitro inhibition of prolyl endopeptidase from 41, No. 9, 1993, pages. 1583-1588.	otidase inhibitors and evaluation of their s n Canine Brain" <u>CHEMICAL AND PHAI</u>	tructure-activity relationships: in RMACEUTICAL BULLETIN., I	
CJ	J. Lin et al.: "Inhibition of depeptidyl peptidepeptidomimetics" PROCEEDINGS OF THE 1998, pages 14020-14024.	ase IV by fluoroolefin-containing n-peptic ENATIONAL ACADEMY OF SCIENCE	dyl-O-hydroxylamine ES OF USA, Vol. 95, November	
CK	KOROM, S., et al "Inhibition of CD26/diper recipients", <u>Transplantation</u> , Vol. 63, 1495 –		gs cardiac allograft survival in ra	
CL	TANKA, S., et al., "Suppression of arthritis 19, No. 1 Pages 15-24, (1997)	TANKA, S., et al., "Suppression of arthritis by the inhibitors of dipeptidyl peptidase IV". <u>Int. J. Immunopharmacol</u> , Vol. 19, No. 1 Pages 15-24, (1997)		
СМ	MENTLEIN, R., et al., "Proteolytic processing of neuropeptide Y and peptide YY by dipeptidyl peptidase IV". Regul. Pept. 49, 133-144 (1993)			
. CN	WETZEL, W., et al., "Effects of the CLIP fragment ACTH 20-24 on the duration of REM sleep episodes". Neuropeptides, 31, 41-45 (1997)			
.co	AMASHEH, S., et al., "Electrophysiological expressed in Xenopus Laevis oocytes". J. Pl		an renal peptide transporter	
СР	DURINX, C.; et al.; "Reference Values for Laboratory Parameters". Clin Chem Lab M			
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FORM PTO-1449 INFORMATION DISCLOSURE STATEMENT		ATTY. DOCKET NO. 20784/6	SERIAL NO. 10/082,001
	•	APPLICANT(S): Demuth et al.	
		FILING DATE: February 22, 2002	ART UNIT: 1646
	OTHER DOCUMENTS (INCLUDING AU	JTHOR, TITLE, DATE, PERTINEN	T PAGES, ETC.)
CQ	GOSSRAU, R.; "Cytochemistry of Membrane	e Proteases". Histochem J, 1985, July;	17 (7):737-71, 1 page.
CR	HAHN, T.; et al.; "Enzyme Histochemical Ev Cultured Villous Explants from Human First page.	ridence for the Presence of Potential Bl Trimester Placentae". Acta Histochem	ood Pressure Regulating Proteases in 1993, December, 95 (2):185-92, 1
CS	HEYMANN, E. et al., "Has Dipeptidyl Peptid 1984, January, 2;62 (1):2-10, 1 page.	dase IV an Effect on Blood Pressure an	d Coagulation." Klin Wochenschr,
СТ	MAGYAR, C.E. et al., "Proximal Tubule Na Hypertension." <u>Am J. Physiol Renal Physiol</u> ,	Transporter Responses are the same du, 2000, August; 279 (2):F358-69, 1 pag	ring Acute and Chronic ge.
CU	PAPIES, B. et al., "Isoenzyme (Lactate Dehydrogenase, Aspartate Aminotransferase) and Dipeptidyl Peptidase IV Activity Changes in Blood Plasma Likely Indicative of Organ Involvement due to Arterial Hypertension." Cor Vasa, 1991; 33 (3):218-26, 1 page.		
CV	QURESHI. N.U.; et al., "Endogenous Neuropeptide Y Mediates Vasoconstriction during Endotoxic and Hemorrhagic Shock". Regul Pept, 1998, September 25; 75-76:215-20, 1 page.		
CW	Index Nominum, International Drug Directory 1992/1993, Medpharm Scientific Publishers, pages 728-729.		
CX	The Merck Index, An Encyclopedia of Chemicals and Drugs, 9th Edition, Merck & Co., Inc., 1976, page 773		
CY	Willms et al., <u>Journal of Clinical Endocrinology Metabolism</u> , "Gastric Emptying, Glucose Responses, and Insulin Secretion after a Liquid Test Meal: Effects of Exogenous Glucagon-Like Peptide-1 (GLP-1)-(7-36) Amide in Type 2 (Noninsulin-Dependent) Diabetic Patients", 1996, 81(1): 327-332.		
CZ	Hoffmann et al., <u>Journal of Chromatography A</u> , "Inhibition of dipeptidyl peptidase IV (DP IV) by anti-DP IV antibodies and non-substrate X-X-Pro- oligopeptides ascertained by capillary eletrophoresis", 1995, 716:355-362.		
CAA	C.B. Welch, <u>Medical Management of Non-Insulin-Dependent (Type II) Diabetes</u> , 3 rd edition, American Diabetes Association, "Diagnosis and Classification" p. 3, 1994, Pharmacologic Intervention (2 pages).		
CAB	Mannucci et al., <u>Diabetes Care</u> , "Effect of Metformin on Glucagon-Like Peptide 1 (GLP-1) and Leptin Levels in Obese Nondiabetic Subjects", 24(3): 489-494, March 2001.		
CAC	Stryer, Biochemistry 3 rd Ed., "Protein Conformation, Dynamics, and Function", 1988, p 191-193.		
CAD	Pauly et al., <u>Regulatory Peptides</u> , "Abstracts Issue: Abstracts from the 11 th International Symposium on Regulatory Peptides", July 15, 1996, 64(1-3): 148 plus cover.		
CAE	Gutniak et al., New England Journal of Med Normal Subjects and Patients With Diabetes	icine, "Antidiabetogenic Effect of Gluc Mellitus", 1992, 326: 1316-1322.	agon-like peptide-1 (7-36) Amide in
CAF	Hendrick et al., Metabolism – Clinical and Ex Rats", January 1993, 42(1): 1-6.	xperimental, "Glucagon-like Peptide-I-	(7-37) Suppresses Hyperglycemia in
Examiner:		Date:	

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FORM PTO-1449 INFORMATION DISCLOSURE STATEMENT		ATTY: DOCKET NO.	SERIAL NO. 10/082,001	
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		APPLICANT(S): Demuth et al.	ADDID	
		FILING DATE: February 22, 2002	ART UNIT: 1646	
	OTHER DOCUMENTS (INCLUDING A	AUTHOR, TITLE, DATE, PERTINENT	PAGES, ETC.)	
CAG	Nauck et al., <u>Diabetologia</u> , "Normalization in Type 2 (non-insulin-dependent) diabetic	of fasting hyperglycaemia by exogenous g patients", (1993), 36: 741-744.	lucagon-like peptide 1 (7-36 amide	
САН	Gutniak et al., <u>Diabetes Care</u> , "Subcutaneou Postprandial Glycemia in NIDDM", Septen	ns Injection of the Incretin Hormone Gluca nber 1994, 17(9): 1039-1044.	gon-Like Peptide 1 Abolishes	
CAI	Deacon et al., <u>Journal of Clinical Endocrino</u> Plasma in Vitro Yields and N-Terminally T 80(3): 952-957.	ology and Metabolism, "Degradation of Gl runcated Peptide That Is a Major Endogen	ucagon-Like Peptide-1 by Human ous Metabolite in Vivo", (1995),	
CAJ	H.A. Smith et al., <u>Veterinary Pathology</u> (for (1972), p 1018-1020.			
CAK	G.G. Duncan, Diseases of Metabolism (Asi	an edition), "Diabetes Mellitus", (1966), p	951-957.	
CAL	T.J. Kieffer et al., "Degradation of Glucose-Dependent Insulinotropic Polypetide and Truncated Glucagon-Like Peptide 1 In Vitro and In Vivo by DP IV", Endocrinology, Vol. 136(8), (1995), p 3585-3596.			
CAM	C.F. Deacon et al., <u>Diabetes</u> , "Both Subcutaneously and Intravenously Administered Glucagon-Like Peptide I Are Rapidly Degraded from the NH ₂ -Terminus in Type II Diabetic Patients and in Healthy Subjects", September 1995, 44: 1126-1131.			
CAN	Pauly et al., Metabolism, "Improved Gluco Inhibitor Ile-Thiazolidide", (1999), 48(3): 3		ptidyl Peptidase IV (CD26)	
CAO	Vidal, (1993), 69th Edition, p. 612-613.			
CAP	Goodman & Gilman's The Pharmacologica	l Basis of Therapeutics, Ninth Edition, (19	996), p. 1510.	
CAQ	Nathan et al., <u>Diabetes Care</u> , "Insulinotropic Action of Glucagonlike Peptide-1-(7-37) in Diabetic and Nondiabetic Subjects", February 1992, 15(2): 270-275.			
CAR	Pschyrembel, Kininisches Wörterbuch 257,	, Auflage, (1994), 9 pages.		
CAS	Frohman et al., <u>Journal of Clin. Invest.</u> , "Rapid Enzymatic Degradation of Growth Hormone-releasing Hormone by Plasma in Vitro and in Vivo to a Biologically Inactive Product Cleaved at the NH ₂ Terminus", Volume 78, October 1986, p 906-913			
CAT	Snow et al., Advances In Medicinal Chemistry, "Boronic Acid Inhibitors of Dipeptidyl Peptidase IV: A New Class of Immunosuppressive Agents", Vol. 3, (1995), p 149-177.			
• CAU	Thorens et al., <u>Diabetes</u> , "Glucagon-Like Pepetide-I and the Control of Insulin Secretion in the Normal State and in NIDDM", (1993), 42:1219-1225.			
CAV	Wakselman et al., "Inhibition of HIV-1 info DPP IV activity of CD26", Abstract P 44 o	ection of CD 26 ⁺ but not CD 26 ⁻ cells by a pf the 24 th European Peptide Symposium, (potent cyclopeptidic inhibitor of th 1996).	
CAW	0.11 7.11 (D) (1.11			
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		APPLICANT(S): Demuth et al.	
		FILING DATE: February 22, 2002	ART UNIT: 1646
	OTHER DOCUMENTS (INCLUDING A	UTHOR, TITLE, DATE, PERTINEN	T PAGES, ETC.)
CAX	Endroczi et al., <u>ACTA PHYSIOL. HUNG.</u> , Thymus-Derived Lymphocytes: Effects of In	"Dipeptidyl peptidase IV (DP IV) and Sonhibitory Pepdides and Zn ²⁺ in Vitro", (1	uperoxide Dismutase Activity in 1990), 75(1): 35-44.
CAY	Lee, H.S. et al., "Cathepsin B Inhibitory Peptides Derived from β-Casein," Peptides 21 (2000) 807-809.		
CAZ	Edwards, J.V. et al., <u>J. Peptide Res.</u> , "Synthe Sequences on Cotton," (1999), 54: 536-543.		erminal Elastase Recognition
СВА	Wettstein, J.G. et al. <u>Pharmacology & Thera</u> (1995), 65(3): 397-414.	ipeutics, "Central Nervous System Pharm	nacology of Neuropeptide Y.",
СВВ	Badia-Elder N.E. et al., <u>Alcoholism Clinical</u> Intake and Anxiety in High and Low Alcoho	and Experimental Research, "Effects of ol Drinking (HAD1/LAD1) Rats", (2000	Neuropeptide Y (NPY) on Ethanol), 24(5): 82A.
СВС	Munglani R. et al., Drugs, Adis Internationa Anxiolytic and Antihypertensive", (1996) 52		Neuropeptide Y Analgesic,

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